THE SCHEDULE

(*See* sections 8 and 16**)**

Standards to be complied with by imported drugs and by drugs manufactured for sale, sold, stocked or exhibited for sale, or distributed.

Class of drug	Standard to be complied with
1. Patent or proprietary medicines. ….	The formula or list of ingredients displayed in the prescribed manner on the label or container.
2. Substances commonly known as vaccines, sera, toxins, toxoids, anti-toxins, and antigens and biological products of such nature. …	The standards maintained at the National Institute for Medical Research, London, and such further standards of strength, quality and purity as may be prescribed.
3. Vitamins, hormones and analogous products.	The standards maintained at the National Institute for Medical Research, London, and such further standards of strength, quality and purity as may be prescribed.
[1][3A. Substances (other than food) intended to affect the structure or any function of the human body or intended to be used for the destruction of vermins or insects which cause disease in human beings or animals. …	Such standards as may be prescribed.]
4. Other drugs …	The standards of identity, purity and strength specified in the latest edition of the British Pharmacopoeia or the British Pharmaceutical Codex or any other prescribed pharmacopoeia, or adopted by the Permanent Commission on Biological Standardisation of the World Health Organization.

[1] Entry No. 3A was inserted by section 16 of the Drugs (Amendment) Act, 1963 (Act No. XXII of 1963).